TRANSMITTAL OF INFORMATION DISCLOSURE STATEMENT Docket No.								
(Under 37 CFR 1.97(b) or 1.97(c)) RLL-293US								
In Re Application Of: Mehta et al. SEP 2 2 2006 in								
Application	Application No. Filing Date Customer No. Group Art Unit Confirmation No.							
10/552,456 08/14/2006 Unknown 26815 1614 2308								
Title: SUBS								
	•		Address to: Commissioner for Paten P.O. Box 1450 Alexandria, VA 22313-14					
			37 CFR 1.97(b)					
of a three appli	The Information Disclosure Statement submitted herewith is being filed within three months of the filing of a national application other than a continued prosecution application under 37 CFR 1.53(d); within three months of the date of entry of the national stage as set forth in 37 CFR 1.491 in an international application; before the mailing of a first Office Action on the merits, or before the mailing of a first Office Action after the filing of a request for continued examination under 37 CFR 1.114.							
			37 CFR 1.97(c)					
CFR Fina	2. The Information Disclosure Statement submitted herewith is being filed after the period specified in 37 CFR 1.97(b), provided that the Information Disclosure Statement is filed before the mailing date of a Final Action under 37 CFR 1.113, a Notice of Allowance under 37 CFR 1.311, or an Action that otherwise closes prosecution in the application, and is accompanied by one of:							
	☐ the statement specified in 37 CFR 1.97(e);							
OR								
☐ the fee set forth in 37 CFR 1.17(p).								
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TRANSMITTAL OF INFORMATION DISCLOSURE STATEMENT (Under 37 CFR 1.97(b) or 1.97(c)) Docket No. RLL-293US							
In Re Applica	ation o	f: Mehta et al.	SEP 2 2 7006 W				
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WAR inclu C I certify account Patent a	ertific that this is beir nd Trac	y credit card. Form F : Information on this in this form. Provide ate of Transmission be document and authorizate gracsimile transmitted lemark Office (Fa Signature Printed Name of Person Signate may only be used unt. Signature	is form may become credit card information of the United States ening Certificate	rmation and Ce I hereby of with the U as first "Commiss 22313-145 Septe	authorization rtificate of Mail certify that this co Inited States Posts class mail in sioner for Patents, 50" [37 CFR 1.8(a)] mber 20, 2006 (Date) Signature of Per Christine	ing by First Class respondence is be all Service with suffi an envelope a P.O. Box 1450, Ale on	ing deposited cient postage ddressed to exandria, VA
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INFORMATION DISCLOSURE CITATION

Docket No.: RLL-293US

Serial No.: 10/552,456

Applicants: MEHTA et al.

Filed: 10/7/2005

Group:

SEP	2	2	2006

/3	U.S. PATENT DOCUMENTS EXAMINER FILING DATE						
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
	A1	3,176,019	3/30/1965	Campbell <i>et al</i> .	260	293.4	
	A2	5,281,601	1/25/1994	Cross et al.	514	320	-u-sgass
	А3	5,397,800	3/14/1995	Alker <i>et al</i> .	514	413	
	A4	5,559,269	9/24/1996	Johansson et al.	564	443	
	A5	5,948,792	9/7/1999	Tsuchiya <i>et al.</i>	514	317	
	A6	6,130,232	10/10/2000	Mase et al.	514	318	
	A7	6,174,900	1/16/2001	Okada <i>et al.</i>	514	317	
			FOREIGN	N PATENT DOCUMENTS	•		
		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION YES NO
	B1	EP 0 325 571	7/26/1989	EPO	C07C	215/54	
	B2	EP 0 388 054	9/19/1990	EPO	C07D	207/08	
	В3	EP 0 801 067	10/15/1997	EPO	C07D	453/02	
	B4	EP 0 823 423	2/11/1998	EPO	C07D	211/46	
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	В6	JP 135958/1994	5/17/1994	Japan	C07D	333/16	
	B7	JP 92921/1994	4/5/1994	Japan	C07C	237/20	
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	В9	WO 93/16018	8/19/1993	PCT	C05F	17/02	
	B10	WO 93/16048	8/19/1993	PCT	C07D	211/26	
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	B12	WO 97/45414	12/4/1997	PCT	C07D	211/58	
	B13	WO 98/00016	1/8/1998	PCT	A01N	43/38	1

DATE CONSIDERED

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ć	B15	WO 98/00132	1/8/1998	PCT		A61K	31/40	
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	B17	WO 98/05641	2/12/1998	PCT		C07D	211/46	
	B18	WO 98/29402	7/9/1998	PCT		C07D	311/20	
		OTHER DOCUME	NTS (Includ	ding Auth	nor, Title, Date, Pe	ertinent P	ages, Etc.)	
	C1	Kubo et al., "Clon muscarinic acetyl	choline rec	eptor", N	ature, 323(2):411	-416 (198	6)	
	C2 Bonner et al., "Identification of a Family of Muscarinic Acetylcholine Receptor Genes", Science, 237:527-531 (1987)							
	Eglen et al., "Muscarinic receptor ligands and their theraputic potential", <i>Current Opinion in Chemical Biology</i> , 3:426-432 (1999)							
	C4	Eglen et al., "Theraputic opportunities from muscarinic receptor research", <i>Trends in Pharmacological Sciences</i> , <u>22</u> (8):409-414 (2001)						
	C5	Felder et al., "Theraputic Opportunities for Muscarinic Receptors in the Central Nervous System", <i>Journal of Medicinal Chemistry</i> , 43(23):4333-4353 (2000)						
	C6 Broadley and Kelly, "Muscarinic Receptor Agonists and Antagonists", <i>Molecules</i> , <u>6</u> :142-193 (2001) C7 Birdsall et al., "Muscarinic receptors: it's a knockout", <i>Trends in Pharmacological Sciences</i> , <u>22</u> (5):215-219 (2001) C8 de Groat and Yoshimura, "Pharmacology of the Lower Urinary Tract", <i>Annual Review of Pharmacology and Toxicology</i> , <u>41</u> :691-721 (2001) C9 Steers, "The future direction of neuro-urology drug research", <i>Current Opinion in CPNS Investigational Drugs</i> , <u>2</u> (3):268-282							
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	C10	Chapple, "Muscarinic receptor antagonists in the treatment of overactive bladder", <i>Urology</i> , 55(Suppl. 5A):33-46 (2000) Stoors, Barret, Wein, "Voiding dysfunction; diagnosis classification and management". In:						
	C11							
	C12	Sagara et al., "Cy M3 Antagonist Di	Sagara et al., "Cyclohexylmethylpiperidinyltriphenylpropioamide: A Selective Muscarinic M3 Antagonist Discriminating against the Other Receptor Subtypes", <i>Journal of Medicinal</i>					
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		Docket No.: RLL-293US	Serial No.: 10/552,456			
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6.04.4		Filed: 10/7/2005	Group:			
SEP 2 2 2006	5 w					
C1 PADE MART	Muscarinic Acetylcholine Receptor (m Submandibular Gland", <i>Life Sciences</i>	Moriya et al., "Affinity Profiles of Various Muscarinic Antagonists for Cloned Human Muscarinic Acetylcholine Receptor (mAChR) Subtypes and mAChRs in Rat Heart and Submandibular Gland", <i>Life Sciences</i> , 64(25):2351-2358 (1999)				
C1	Cheng and Prusoff, "Relationship between the inhibition constant (<i>K1</i>) and the concentration of inhibitor which causes 50 per cent inhibition (<i>I50</i>) of an enzymatic reaction", <i>Biochemical Pharmacology</i> , 22:3099-3108 (1973)					
C1	Receptor Antagonist, via Copper-Ass	Andersson et al, "Asymmetric Total Synthesis of (+)-Tolterodine, a New Muscarinic Receptor Antagonist, via Copper-Assisted Asymmetric Conjugate Addition of Aryl Gridngard Reagents to 3-Phenyl-prop-2-enoyl-oxazolidinones", <i>Journal of Organic</i>				

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